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60/566,867 30 April 2004 (30.04.2004) (71) Applicant (for all designated States except US): TI-BOTEC PHARMACEUTICALS LTD. [IE/IE]; Little

(72) Inventors; and

- BONFANTI, (75) Inventors/Applicants (for US only): Jean-François [FR/FR]; 4 bis Route Nationale, F-27430 Andé (FR). ANDRIES, Koenraad, Jozef, Lodewijk [BE/BE]; Oosteneinde 9, B-2340 Beerse (BE). FORTIN, Jérôme, Michel, Claude [FR/FR]; 22, rue des acacias, F-27460 Igoville (FR). MULLER, Philippe [FR/FR]; 17 Route d'Herqueville, F-27430 Andé (FR). DOU-BLET, Frédéric, Marc, Maurice [FR/FR]; 1646, Route de Neufchatel, F-76230 Isneauville (FR). MEYER, Christophe [FR/FR]; 428 rue de Gouy, F-76520 Les Authieux s/l Port St Ouen (FR). WILLEBRORDS, Rudy, Edmond [BE/BE]; Vaartstraat 70, B-2330 Merksplas (BE). GEVERS, Tom, Valerius, Josepha [BE/BE]; Burgemeester Bossaertlaan 17, B-2350 Vosselaar (BE). TIMMERMAN, Philip, Maria, Martha, Bern [BE/BE]; Vijversstraat 211, B-3550 Hasselt (BE).
- (74) Agent: WANTE, Dirk; Tibotec-Virco Comm. VA, Generaal De Wittelaan L 11B 3, B-2800 Mechelen (BE).

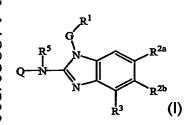
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Declarations under Rule 4.17:

- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii)) for the following designations AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VC, VN, YU, ZA, ZM, ZW, ARIPO patent (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG)
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii)) for all designations

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(54) Title: 5- OR 6-SUBSTITUTED BENZIMIDAZOLE DERIVATIVES AS INHIBITORS OF RESPIRATORY SYNCYTIAL VIRUS REPLICATION



(57) Abstract: The present invention concerns 5- or 6-substituted-benzimidazole derivatives having inhibitory activity on the replication of the respiratory syncytial virus and having the formula (I) a prodrug, N-oxide, addition salt, quaternary amine, metal complex or stereochemically isomeric form thereof wherein Q is Ar2, R6, pyrrolidinyl substituted with R⁶, piperidinyl substituted with R⁶ or homopiperidinyl substituted with R⁶, G is a direct bond or optionally substituted C₁₋₁₀alkanediyl; R¹ is Ar¹ or a monocyclic or bicyclic heterocycle; one of R ^{2a} and R^{2b} is cyanoC_{1.6}alkyl, cyanoC_{2.6}alkenyl, Ar³C_{1.6}alkyl, Het- $-C_{1\text{-}6}alkyl,\ N(R^{8a}R^{8b})C_{1\text{-}6}alkyl,\ Ar^3C_{2\text{-}6}alkenyl,\ Het-C_{2\text{-}6}alkenyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Het-C_{2\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Het-C_{2\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Het-C_{2\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Het-C_{2\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Het-C_{2\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Het-C_{2\text{-}6}alkyl,\ Ar^3aminoC_{1\text{-}6}alkyl,\ Ar^$ aminoC₁₋₆alkyl, Ar³thioC₁₋₆ alkyl, Het-thioC₁₋₆alkyl, Ar³sulfonylC₁₋₆ alkyl, HetsulfonylC₁.

6alkyl, Ar3aminocarbonyl, Het-aminocarbonyl, Ar3(CH2)naminocarbonyl, Het-(CH2)naminocarbonyl, Ar3carbonylamino, Het-carbonylamino, Ar3(CH₂)_ncarbonylamino, Het-(CH₂)_ncarbonylamino, and the other one of R^{2a} and R^{2b} is hydrogen; in case R^{2a} is hydrogen, then R3 is hydrogen; in case R2b is hydrogen, the R3 is hydrogen or C1-6alkyl. It further concerns their preparation and compositions comprising them, as well as their use as a medicine.



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- of inventorship (Rule 4.17(iv)) for US only

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